

(FILE 'HOME' ENTERED AT 13:42:39 ON 09 NOV 2006)

FILE 'REGISTRY' ENTERED AT 13:42:51 ON 09 NOV 2006

L1           STRUCTURE UPLOADED  
L2           0 S L1  
L3           23 S L1 SSS FULL  
L4           STRUCTURE UPLOADED  
L5           1 S L4  
L6           33 S L4 SSS FULL

FILE 'CAPLUS' ENTERED AT 13:45:49 ON 09 NOV 2006

L7           0 S L3 AND L6  
L8           0 S L6 AND (PDEV OR PHOSPHODIESTERASE(W) (V OR 5))  
L9           0 S L3 AND ANTICHOLINERGIC  
L10          1 S L6 AND (PDEIV OR PDE4 OR PHOSPHODIESTERASE(W) (IV OR 4))

FILE 'REGISTRY' ENTERED AT 13:47:56 ON 09 NOV 2006

L11          3 S ENPROFYLLINE/CN OR THEOPHYLLINE/CN OR ROFLUMILAST/CN OR (BAY-

FILE 'CAPLUS' ENTERED AT 13:48:43 ON 09 NOV 2006

L12          1 S L6 AND L11  
L13          5 S L3  
L14          26 S L6  
L15          0 S L14 NOT PY>2002  
L16          0 S L14 NOT PY>2003  
L17          1 S L14 NOT PY>2004

=> file registry  
COST IN U.S. DOLLARS

SINCE FILE ENTRY	TOTAL SESSION
0.21	0.21

FULL ESTIMATED COST

FILE 'REGISTRY' ENTERED AT 13:42:51 ON 09 NOV 2006  
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STRUCTURE FILE UPDATES: 8 NOV 2006 HIGHEST RN 912757-80-3  
DICTIONARY FILE UPDATES: 8 NOV 2006 HIGHEST RN 912757-80-3

New CAS Information Use Policies, enter HELP USAGETERMS for details.

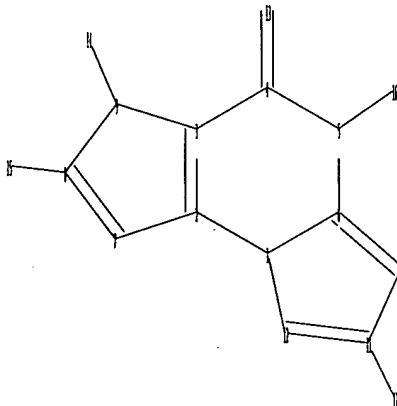
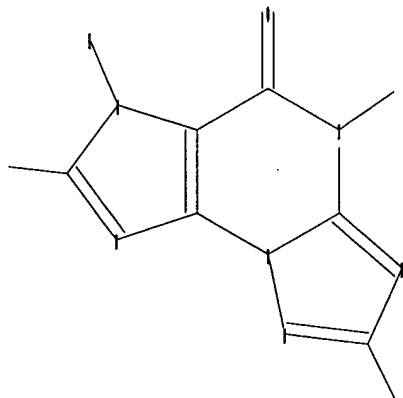
TSCA INFORMATION NOW CURRENT THROUGH June 30, 2006

Please note that search-term pricing does apply when  
conducting SmartSELECT searches.

REGISTRY includes numerically searchable data for experimental and  
predicted properties as well as tags indicating availability of  
experimental property data in the original document. For information  
on property searching in REGISTRY, refer to:

<http://www.cas.org/ONLINE/UG/regprops.html>

=>  
Uploading C:\Program Files\Stnexp\Queries\10613783pde4.str



chain nodes :

13 14 15 16 17

ring nodes :

1 2 3 4 5 6 7 8 9 10 11 12

chain bonds :

4-13 5-16 8-15 9-14 11-17

ring bonds :

1-2 1-6 1-12 2-3 2-7 3-4 3-9 4-5 5-6 6-10 7-8 8-9 10-11 11-12

exact/norm bonds :

1-2 1-6 1-12 2-3 2-7 3-4 3-9 4-5 4-13 5-6 5-16 6-10 7-8 8-9 10-11 11-12

exact bonds :

8-15 9-14 11-17

Match level :

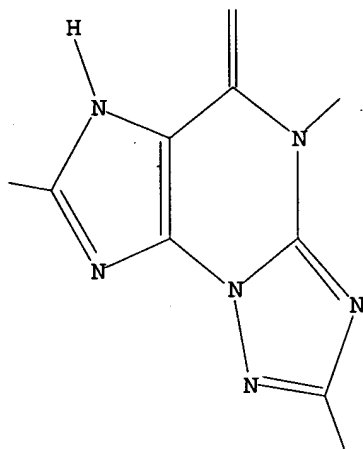
1:Atom 2:Atom 3:Atom 4:Atom 5:Atom 6:Atom 7:Atom 8:Atom 9:Atom 10:Atom  
11:Atom 12:Atom 13:CLASS 14:CLASS 15:CLASS 16:CLASS 17:CLASS

L1 STRUCTURE UPLOADED

=> d l1

L1 HAS NO ANSWERS

L1 STR



Structure attributes must be viewed using STN Express query preparation.

=> s l1

SAMPLE SEARCH INITIATED 13:43:30 FILE 'REGISTRY'

SAMPLE SCREEN SEARCH COMPLETED - 6 TO ITERATE

100.0% PROCESSED 6 ITERATIONS

0 ANSWERS

SEARCH TIME: 00.00.01

FULL FILE PROJECTIONS: ONLINE \*\*COMPLETE\*\*

BATCH \*\*COMPLETE\*\*

PROJECTED ITERATIONS: 6 TO 266

PROJECTED ANSWERS: 0 TO 0

L2 0 SEA SSS SAM L1

=> s l1 sss full

FULL SEARCH INITIATED 13:43:38 FILE 'REGISTRY'

FULL SCREEN SEARCH COMPLETED - 174 TO ITERATE

100.0% PROCESSED 174 ITERATIONS

23 ANSWERS

SEARCH TIME: 00.00.01

L3 23 SEA SSS FUL L1

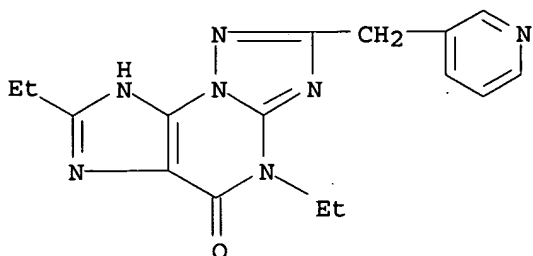
=> d l23 scan

L23 NOT FOUND

The L-number has not been used in the current session or has been deleted.

=> d 13 scan

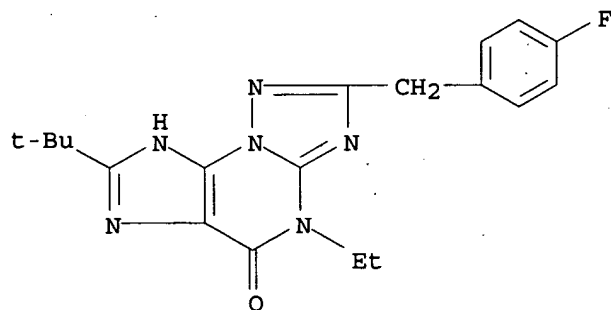
L3 23 ANSWERS REGISTRY COPYRIGHT 2006 ACS on STN  
IN 4H-[1,2,4]Triazolo[5,1-b]purin-5(6H)-one, 4,7-diethyl-2-(3-pyridinylmethyl)- (9CI)  
MF C16 H17 N7 O



\*\*PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT\*\*

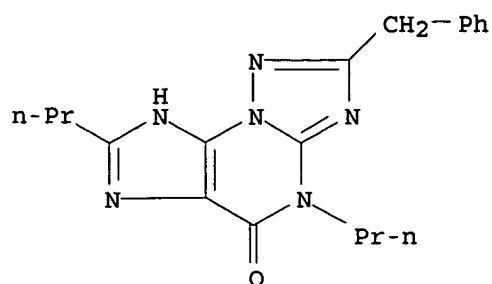
HOW MANY MORE ANSWERS DO YOU WISH TO SCAN? (1):4

L3 23 ANSWERS REGISTRY COPYRIGHT 2006 ACS on STN  
IN 4H-[1,2,4]Triazolo[5,1-b]purin-5(6H)-one, 7-(1,1-dimethylethyl)-4-ethyl-2-[(4-fluorophenyl)methyl]- (9CI)  
MF C19 H21 F N6 O



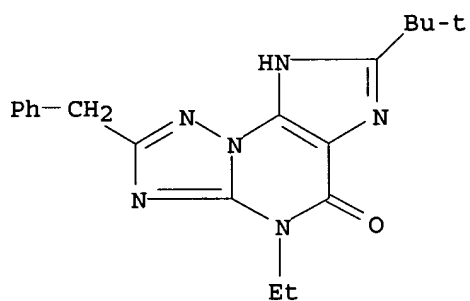
\*\*PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT\*\*

L3 23 ANSWERS REGISTRY COPYRIGHT 2006 ACS on STN  
IN 4H-[1,2,4]Triazolo[5,1-b]purin-5(6H)-one, 2-(phenylmethyl)-4,7-dipropyl- (9CI)  
MF C19 H22 N6 O



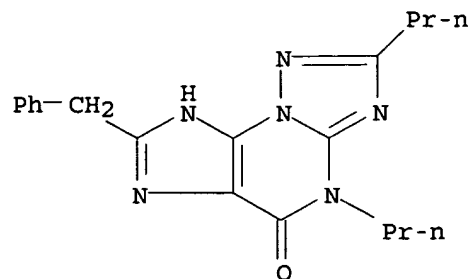
\*\*PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT\*\*

L3 23 ANSWERS REGISTRY COPYRIGHT 2006 ACS on STN  
 IN 4H-[1,2,4]Triazolo[5,1-b]purin-5(6H)-one, 7-(1,1-dimethylethyl)-4-ethyl-2-(phenylmethyl)- (9CI)  
 MF C19 H22 N6 O



\*\*PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT\*\*

L3 23 ANSWERS REGISTRY COPYRIGHT 2006 ACS on STN  
 IN 4H-[1,2,4]Triazolo[5,1-b]purin-5(6H)-one, 7-(phenylmethyl)-2,4-dipropyl- (9CI)  
 MF C19 H22 N6 O



\*\*PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT\*\*

HOW MANY MORE ANSWERS DO YOU WISH TO SCAN? (1):

Uploading

'UPLOAD SSTN' IS NOT VALID HERE

To display more answers, enter the number of answers you would like to see. To end the display, enter "NONE", "N", "0", or "END".

HOW MANY MORE ANSWERS DO YOU WISH TO SCAN? (1):C:\Program

Files\Stnexp\Queries\10614365anticholinergic.str

YOU WISH TO SCAN? (1):

'0 SZ' @-#&1~" J\*' IS NOT VALID HERE

To display more answers, enter the number of answers you would like to see. To end the display, enter "NONE", "N", "0", or "END".

HOW MANY MORE ANSWERS DO YOU WISH TO SCAN? (1):

'0 SZ' @-#&1~" J\*' IS NOT VALID HERE

To display more answers, enter the number of answers you would like to see. To end the display, enter "NONE", "N", "0", or "END".

HOW MANY MORE ANSWERS DO YOU WISH TO SCAN? (1):

'0 SZ' @-#&1~" J\*' IS NOT VALID HERE

To display more answers, enter the number of answers you would like to see. To end the display, enter "NONE", "N", "0", or "END".

HOW MANY MORE ANSWERS DO YOU WISH TO SCAN? (1):

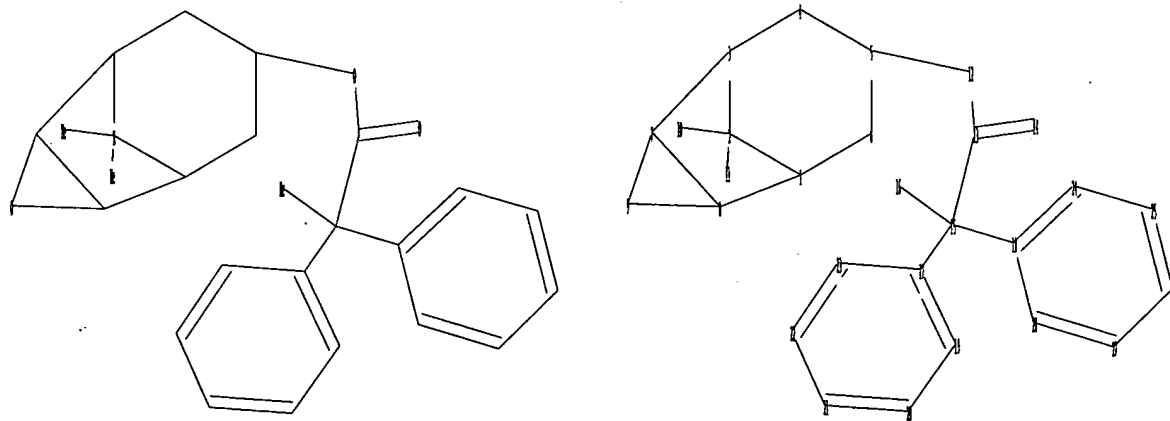
'0 SZ' @-#&1~" J\*' IS NOT VALID HERE

To display more answers, enter the number of answers you would like to see. To end the display, enter "NONE", "N", "0", or "END".

HOW MANY MORE ANSWERS DO YOU WISH TO SCAN? (1):0

=>

Uploading C:\Program Files\Stnexp\Queries\10614365anticholinergic.str



chain nodes :

10 11 12 13 14 15 16

ring nodes :

1 2 3 4 5 6 7 8 9 17 18 19 20 21 22 23 24 25 26 27 28

chain bonds :

2-10 2-11 5-12 12-13 13-14 13-15 15-16 15-17 15-18

ring bonds :

1-2 1-6 1-8 2-3 3-4 3-7 4-5 5-6 7-8 7-9 8-9 17-19 17-23 18-24 18-28  
19-20 20-21 21-22 22-23 24-25 25-26 26-27 27-28

exact/norm bonds :

1-2 1-6 1-8 2-3 3-4 3-7 4-5 5-6 5-12 7-8 7-9 8-9 12-13 13-14

exact bonds :

2-10 2-11 13-15 15-16 15-17 15-18

normalized bonds :

17-19 17-23 18-24 18-28 19-20 20-21 21-22 22-23 24-25 25-26 26-27 27-28

Match level :

1:Atom 2:Atom 3:Atom 4:Atom 5:Atom 6:Atom 7:Atom 8:Atom 9:Atom 10:CLASS

11:CLASS 12:CLASS 13:CLASS 14:CLASS 15:CLASS 16:CLASS 17:Atom 18:Atom

19:Atom 20:Atom 21:Atom

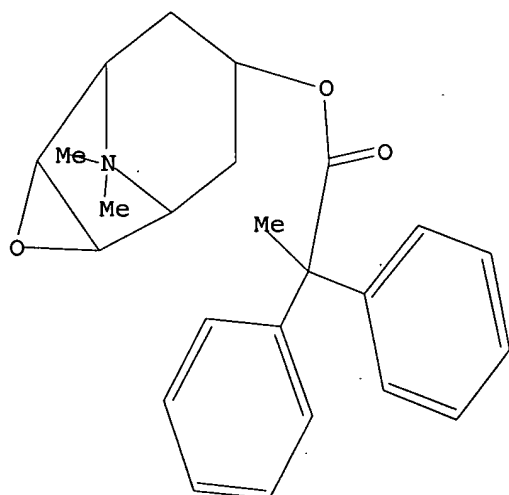
22:Atom 23:Atom 24:Atom 25:Atom 26:Atom 27:Atom 28:Atom

L4 STRUCTURE UPLOADED

=> d l4

L4 HAS NO ANSWERS

L4 STR



Structure attributes must be viewed using STN Express query preparation.

=> s l4

SAMPLE SEARCH INITIATED 13:45:08 FILE 'REGISTRY'

SAMPLE SCREEN SEARCH COMPLETED - 2 TO ITERATE

100.0% PROCESSED 2 ITERATIONS

1 ANSWERS

SEARCH TIME: 00.00.01

FULL FILE PROJECTIONS: ONLINE \*\*COMPLETE\*\*  
BATCH \*\*COMPLETE\*\*

PROJECTED ITERATIONS: 2 TO 124

PROJECTED ANSWERS: 1 TO 80

L5 1 SEA SSS SAM L4

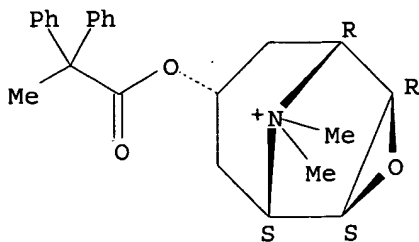
=> d l5 scan

L5 1 ANSWERS REGISTRY COPYRIGHT 2006 ACS on STN

IN 3-Oxa-9-azoniatricyclo[3.3.1.0<sup>2,4</sup>]nonane, 9,9-dimethyl-7-(1-oxo-2,2-diphenylpropoxy)-, (1 $\alpha$ ,2 $\beta$ ,4 $\beta$ ,5 $\alpha$ ,7 $\beta$ )-, (2E)-2-butenedioate (1:1) (9CI)  
 MF C24 H28 N O3 . C4 H3 O4

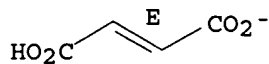
CM 1

Relative stereochemistry.



CM 2

Double bond geometry as shown.



ALL ANSWERS HAVE BEEN SCANNED

=> s l4 sss full  
 FULL SEARCH INITIATED 13:45:22 FILE 'REGISTRY'  
 FULL SCREEN SEARCH COMPLETED - 38 TO ITERATE

100.0% PROCESSED 38 ITERATIONS 33 ANSWERS  
 SEARCH TIME: 00.00.01

L6 33 SEA SSS FUL L4

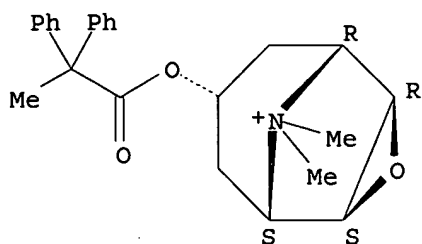
=> d l6 scan

L6 33 ANSWERS REGISTRY COPYRIGHT 2006 ACS on STN  
 IN 3-Oxa-9-azoniatricyclo[3.3.1.0<sup>2,4</sup>]nonane, 9,9-dimethyl-7-(1-oxo-2,2-diphenylpropoxy)-, bromide, (1 $\alpha$ ,2 $\beta$ ,4 $\beta$ ,5 $\alpha$ ,7 $\beta$ )-, mixt. with rel-N-[2-hydroxy-5-[(1R)-1-hydroxy-2-[[ (1R)-2-(4-methoxyphenyl)-1-methylethyl]amino]ethyl]phenyl]formamide (2E)-2-butenedioate (2:1) (salt) dihydrate (9CI)  
 MF C24 H28 N O3 . C19 H24 N2 O4 . 1/2 C4 H4 O4 . Br . H2 O  
 CI MXS

CM 1

Relative stereochemistry.



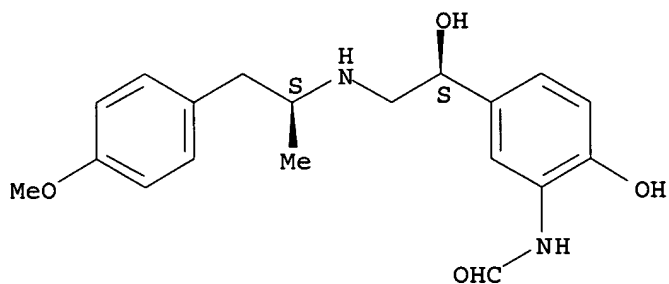


● Br<sup>-</sup>

CM 2

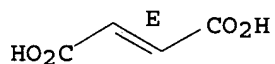
CM 3

Relative stereochemistry.



CM 4

Double bond geometry as shown.



HOW MANY MORE ANSWERS DO YOU WISH TO SCAN? (1):4

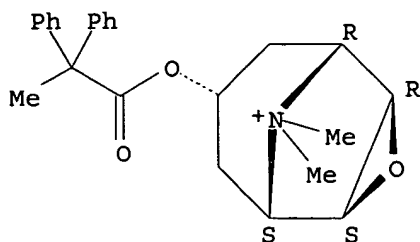
L6 33 ANSWERS REGISTRY COPYRIGHT 2006 ACS on STN  
 IN 3-Oxa-9-azoniatricyclo[3.3.1.0<sup>2,4</sup>]nonane, 9,9-dimethyl-7-(1-oxo-2,2-diphenylpropoxy)-, bromide, (1 $\alpha$ ,2 $\beta$ ,4 $\beta$ ,5 $\alpha$ ,7 $\beta$ )-, mixt. with N-[2-hydroxy-5-[(1R)-1-hydroxy-2-[[[(1R)-2-(4-methoxyphenyl)-1-methylethyl]amino]ethyl]phenyl]formamide (2E)-2-butenedioate (2:1) (salt) dihydrate (9CI)

MF C24 H28 N O3 . C19 H24 N2 O4 . 1/2 C4 H4 O4 . Br . H2 O

CI MXS

CM 1

Relative stereochemistry.

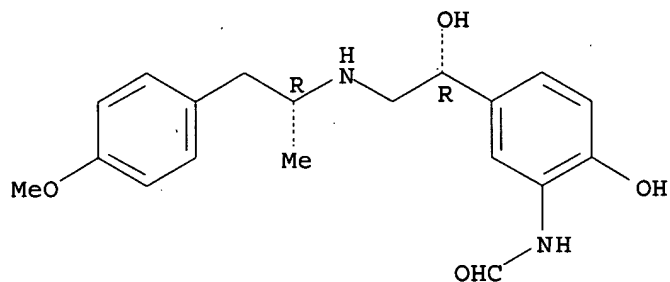


● Br<sup>-</sup>

CM 2

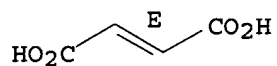
CM 3

Absolute stereochemistry. Rotation (-).



CM 4

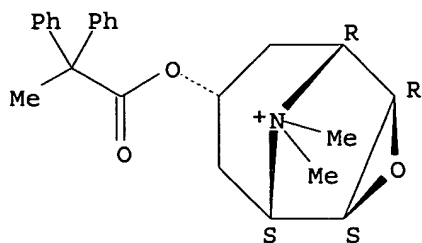
Double bond geometry as shown.



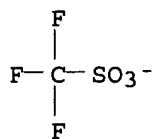
L6 33 ANSWERS REGISTRY COPYRIGHT 2006 ACS on STN  
 IN 3-Oxa-9-azoniatricyclo[3.3.1.0.2,4]nonane, 9,9-dimethyl-7-(1-oxo-2,2-diphenylpropoxy)-, (1 $\alpha$ ,2 $\beta$ ,4 $\beta$ ,5 $\alpha$ ,7 $\beta$ )-, salt with trifluoromethanesulfonic acid (1:1) (9CI)  
 MF C24 H28 N O3 . C F3 O3 S

CM 1

Relative stereochemistry.



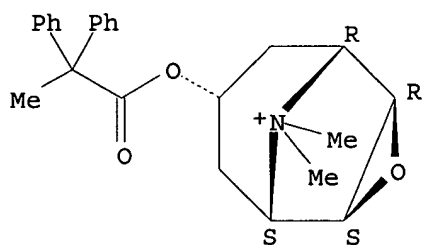
CM 2



L6 33 ANSWERS REGISTRY COPYRIGHT 2006 ACS on STN  
 IN 3-Oxa-9-azoniatricyclo[3.3.1.0<sup>2,4</sup>]nonane, 9,9-dimethyl-7-(1-oxo-2,2-diphenylpropoxy)-, bromide, (1 $\alpha$ ,2 $\beta$ ,4 $\beta$ ,5 $\alpha$ ,7 $\beta$ )-, mixt. with cis-4-cyano-4-[3-(cyclopentyloxy)-4-methoxyphenyl]cyclohexanecarboxylic acid (9CI)  
 MF C24 H28 N O3 . C20 H25 N O4 . Br  
 CI MXS

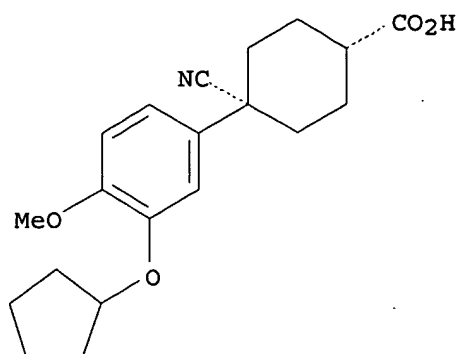
CM 1

Relative stereochemistry.



CM 2

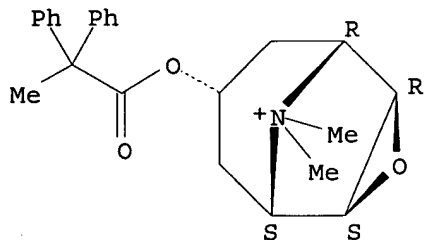
Relative stereochemistry.



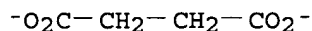
L6 33 ANSWERS REGISTRY COPYRIGHT 2006 ACS on STN  
 IN 3-Oxa-9-azoniatricyclo[3.3.1.0<sup>2,4</sup>]nonane, 9,9-dimethyl-7-(1-oxo-2,2-diphenylpropoxy)-, (1 $\alpha$ ,2 $\beta$ ,4 $\beta$ ,5 $\alpha$ ,7 $\beta$ )-, butanedioate (2:1) (9CI)  
 MF C24 H28 N O3 . 1/2 C4 H4 O4

CM 1

Relative stereochemistry.



CM 2



HOW MANY MORE ANSWERS DO YOU WISH TO SCAN? (1):0

=> file caplus

COST IN U.S. DOLLARS

SINCE FILE	TOTAL
ENTRY	SESSION
335.20	335.41

FULL ESTIMATED COST

FILE 'CAPLUS' ENTERED AT 13:45:49 ON 09 NOV 2006

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FILE COVERS 1907 - 9 Nov 2006 VOL 145 ISS 20  
FILE LAST UPDATED: 8 Nov 2006 (20061108/ED)

Effective October 17, 2005, revised CAS Information Use Policies apply. They are available for your review at:

<http://www.cas.org/infopolicy.html>

=> s l3 and l6

5 L3

26 L6

L7 0 L3 AND L6

=> s l6 and (PDEV or phosphodiesterase(w) (V or 5))

26 L6

14 PDEV

26215 PHOSPHODIESTERASE

1087904 V

6240147 5

1068 PHOSPHODIESTERASE(W) (V OR 5)

L8 0 L6 AND (PDEV OR PHOSPHODIESTERASE(W) (V OR 5))

=> s l3 and anticholinergic

5 L3

5118 ANTICHOLINERGIC

L9 0 L3 AND ANTICHOLINERGIC

=> s l6 and (PDEIV or PDE4 or phosphodiesterase(w) (IV or 4))

26 L6

24 PDEIV

1193 PDE4

26215 PHOSPHODIESTERASE

520751 IV

5451214 4

1494 PHOSPHODIESTERASE(W) (IV OR 4)

L10 1 L6 AND (PDEIV OR PDE4 OR PHOSPHODIESTERASE(W) (IV OR 4))

=> d l10 ti abs bib

L10 ANSWER 1 OF 1 CAPLUS COPYRIGHT 2006 ACS on STN

TI Pharmaceutical compositions comprising anticholinergic agents and phosphodiesterase IV (PDE-IV) inhibitors for the treatment of respiratory diseases

AB The invention provides pharmaceutical compns. comprising anticholinergic agents and PDE-IV inhibitors, as well as a method for the production and use thereof in the treatment of respiratory diseases. Powder inhalant formulations are included.

AN 2004:41257 CAPLUS <<LOGINID::20061109>>

DN 140:87709

TI Pharmaceutical compositions comprising anticholinergic agents and phosphodiesterase IV (PDE-IV) inhibitors for the treatment of respiratory diseases

IN Pairat, Michel; Meade, Christopher John Montague; Pieper, Michael P.

PA Boehringer Ingelheim Pharma G.m.b.H. & Co. K.-G., Germany

SO PCT Int. Appl., 37 pp.

CODEN: PIXXD2

DT Patent

LA German  
FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2004004704	A1	20040115	WO 2003-EP6668	20030625
	W:	AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW			
	RW:	GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG			
	DE 10230769	A1	20040122	DE 2002-10230769	20020709
	CA 2492026	AA	20040115	CA 2003-2492026	20030625
	AU 2003242755	A1	20040123	AU 2003-242755	20030625
	EP 1521576	A1	20050413	EP 2003-762509	20030625
	R:	AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR, BG, CZ, EE, HU, SK			
	JP 2005532379	T2	20051027	JP 2004-518566	20030625
	US 2004058950	A1	20040325	US 2003-614365	20030707
PRAI	DE 2002-10230769	A	20020709		
	US 2002-407895P	P	20020903		
	WO 2003-EP6668	W	20030625		
OS	MARPAT 140:87709				

RE.CNT 9 THERE ARE 9 CITED REFERENCES AVAILABLE FOR THIS RECORD  
ALL CITATIONS AVAILABLE IN THE RE FORMAT

=> file registry

COST IN U.S. DOLLARS

SINCE FILE	TOTAL
ENTRY	SESSION
24.08	359.49

FULL ESTIMATED COST

DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS)

SINCE FILE	TOTAL
ENTRY	SESSION
-0.75	-0.75

CA SUBSCRIBER PRICE

FILE 'REGISTRY' ENTERED AT 13:47:56 ON 09 NOV 2006  
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provided by InfoChem.

STRUCTURE FILE UPDATES: 8 NOV 2006 HIGHEST RN 912757-80-3  
DICTIONARY FILE UPDATES: 8 NOV 2006 HIGHEST RN 912757-80-3

New CAS Information Use Policies, enter HELP USAGETERMS for details.

TSCA INFORMATION NOW CURRENT THROUGH June 30, 2006

Please note that search-term pricing does apply when  
conducting SmartSELECT searches.

REGISTRY includes numerically searchable data for experimental and  
predicted properties as well as tags indicating availability of  
experimental property data in the original document. For information  
on property searching in REGISTRY, refer to:

<http://www.cas.org/ONLINE/UG/regprops.html>

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      1 THEOPHYLLINE/CN
      1 ROFLUMILAST/CN
      0 (BAY-198004)/CN
L11      3 ENPROFYLLINE/CN OR THEOPHYLLINE/CN OR ROFLUMILAST/CN OR (BAY-198
      004)/CN
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                               ENTRY          SESSION
FULL ESTIMATED COST          19.48          378.97

DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS)  SINCE FILE          TOTAL
                                               ENTRY          SESSION
CA SUBSCRIBER PRICE          0.00          -0.75
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 FILE LAST UPDATED: 8 Nov 2006 (20061108/ED)

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=> s l6 and L11
      26 L6
      14693 L11
L12      1 L6 AND L11
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=> d l12 ti abs bib
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L12 ANSWER 1 OF 1 CAPLUS COPYRIGHT 2006 ACS on STN
TI Pharmaceutical compositions comprising anticholinergic agents and
phosphodiesterase IV (PDE-IV) inhibitors for the treatment of respiratory
diseases
AB The invention provides pharmaceutical compns. comprising anticholinergic
agents and PDE-IV inhibitors, as well as a method for the production and use
thereof in the treatment of respiratory diseases. Powder inhalant
formulations are included.
AN 2004:41257 CAPLUS <<LOGINID::20061109>>
DN 140:87709
TI Pharmaceutical compositions comprising anticholinergic agents and
phosphodiesterase IV (PDE-IV) inhibitors for the treatment of respiratory
diseases
IN Pairat, Michel; Meade, Christopher John Montague; Pieper, Michael P.
PA Boehringer Ingelheim Pharma G.m.b.H. & Co. K.-G., Germany
```

SO PCT Int. Appl., 37 pp.

CODEN: PIXXD2

DT Patent

LA German

FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2004004704	A1	20040115	WO 2003-EP6668	20030625
	W:			AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW	
	RW:			GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG	
	DE 10230769	A1	20040122	DE 2002-10230769	20020709
	CA 2492026	AA	20040115	CA 2003-2492026	20030625
	AU 2003242755	A1	20040123	AU 2003-242755	20030625
	EP 1521576	A1	20050413	EP 2003-762509	20030625
	R:			AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR, BG, CZ, EE, HU, SK	
	JP 2005532379	T2	20051027	JP 2004-518566	20030625
	US 2004058950	A1	20040325	US 2003-614365	20030707
PRAI	DE 2002-10230769	A	20020709		
	US 2002-407895P	P	20020903		
	WO 2003-EP6668	W	20030625		

OS MARPAT 140:87709

RE.CNT 9 THERE ARE 9 CITED REFERENCES AVAILABLE FOR THIS RECORD  
ALL CITATIONS AVAILABLE IN THE RE FORMAT

=> s 13

L13 5 L3

=> d l13 1-5 ti

L13 ANSWER 1 OF 5 CAPLUS COPYRIGHT 2006 ACS on STN

TI Preparation of imidazotriazolopyrimidines as adenosine receptor antagonists

L13 ANSWER 2 OF 5 CAPLUS COPYRIGHT 2006 ACS on STN

TI Inhalant compositions containing anticholinergics and PDE IV inhibitors

L13 ANSWER 3 OF 5 CAPLUS COPYRIGHT 2006 ACS on STN

TI Tricyclic nitrogen heterocycles as phosphodiesterase IV inhibitors

L13 ANSWER 4 OF 5 CAPLUS COPYRIGHT 2006 ACS on STN

TI Preparation of imidazotriazolopyrimidines as adenosine receptor antagonists

L13 ANSWER 5 OF 5 CAPLUS COPYRIGHT 2006 ACS on STN

TI Imidazotriazolopyrimidines as adenosine antagonists

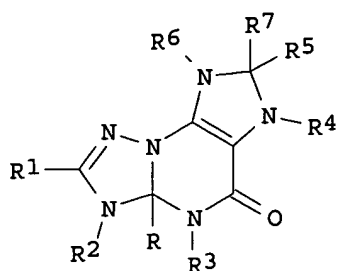
=> d l13 1 2 3 4 5 ti abs bib

L13 ANSWER 1 OF 5 CAPLUS COPYRIGHT 2006 ACS on STN

TI Preparation of imidazotriazolopyrimidines as adenosine receptor antagonists

GI





I

AB Title compds. [I; R1 = H, alkyl, phenyl(alkyl), alkoxy carbonyl, etc.; R2 or R3 = alkyl, alkenyl, benzyl; RR2 or RR3 = bond; R4 or R6 = H, alkyl(amino), CH2Ph, etc.; R4R7 or R6R7 = bond; R5 = H, alkyl, phenyl(alkyl), etc.] were prepared Thus, 7-amino-2-[(4-methoxybenzyloxy)methyl]-s-triazolo[1,5-a]pyrimidine-5-one was converted in 10 steps to I (RR2 = bond, R1 = CH2OPh, R3 = Et, R4 or R6 = H, R4R7 or R6R7 = bond, R5 = cyclopentyl). Data for biol. activity of I were given.

AN 2002:942787 CAPLUS <<LOGINID::20061109>>

DN 138:14073

TI Preparation of imidazotriazolopyrimidines as adenosine receptor antagonists

IN Blech, Stefan; Carter, Adrian; Gaida, Wolfram; Hoffmann, Matthias; Kuefner-Muehl, Ulrike; Meade, Christopher John Montague; Pohl, Gerald; Kummer, Werner; Lehr, Erich; Mierau, Joachim; Weiser, Thomas

PA Boehringer Ingelheim Pharma KG, Germany

SO U.S., 34 pp., Cont.-in-part of U.S. Ser. No. 333,621, abandoned.

CODEN: USXXAM

DT Patent

LA English

FAN.CNT 3

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	US 6492377	B1	20021210	US 2000-559806	20000426
	WO 2000012511	A1	20000309	WO 1998-EP5455	19980827
	W: AU, BG, BR, BY, CA, CN, CZ, EE, HU, ID, IL, JP, KR, KZ, LT, LV, MX, NO, NZ, PL, RO, RU, SG, SI, SK, TR, UA, US, UZ, VN, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM				
	RW: AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE				
	ZA 9808189	A	20000110	ZA 1998-8189	19980908
	BR 9900187	A	20000502	BR 1999-187	19990127
	MX 9905843	A	20000331	MX 1999-5843	19990621
PRAI	US 1998-90586P	P	19980625		
	US 1998-90587P	P	19980625		
	WO 1998-EP5455	A2	19980827		
	US 1999-333408	A2	19990615		
	US 1999-333621	B2	19990615		

OS MARPAT 138:14073

RE.CNT 10 THERE ARE 10 CITED REFERENCES AVAILABLE FOR THIS RECORD

ALL CITATIONS AVAILABLE IN THE RE FORMAT

L13 ANSWER 2 OF 5 CAPLUS COPYRIGHT 2006 ACS on STN

TI Inhalant compositions containing anticholinergics and PDE IV inhibitors

AB The invention relates to drug compns. based on anticholinergics and PDE IV inhibitors, to methods for their production, and to their use as inhalants for the treatment of respiratory tract diseases. Thus an inhalation powder was composed of capsules that contained (µg/capsule): tiotropium

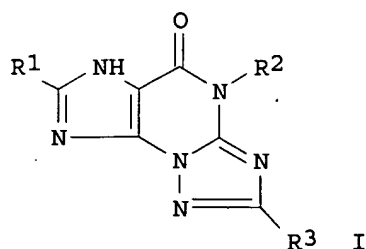
bromide 21.7; AWD-12-281 200; lactose 4778.3.

AN 2002:695761 CAPLUS <<LOGINID::20061109>>  
DN 137:237718  
TI Inhalant compositions containing anticholinergics and PDE IV inhibitors  
IN Meade, Christopher John Montague; Pairet, Michel; Pieper, Michael Paul  
PA Boehringer Ingelheim Pharma K.-G., Germany  
SO PCT Int. Appl., 34 pp.  
CODEN: PIXXD2  
DT Patent  
LA German  
FAN.CNT 14

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2002069945	A2	20020912	WO 2002-EP1988	20020226
	WO 2002069945	A3	20030130		
	W:	AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZM, ZW			
	RW:	GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG			
	DE 10110772	A1	20020912	DE 2001-10110772	20010307
	CA 2439763	AA	20020912	CA 2002-2439763	20020226
	EP 1372649	A2	20040102	EP 2002-727329	20020226
	R:	AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR			
	JP 2004521134	T2	20040715	JP 2002-569122	20020226
	BR 2002007883	A	20040727	BR 2002-7883	20020226
	NZ 528621	A	20050429	NZ 2002-528621	20020226
	CN 1649588	A	20050803	CN 2002-805346	20020226
	ZA 2003006221	A	20040722	ZA 2003-6221	20030812
PRAI	DE 2001-10110772	A	20010307		
	WO 2002-EP1988	W	20020226		
OS	MARPAT 137:237718				

L13 ANSWER 3 OF 5 CAPLUS COPYRIGHT 2006 ACS on STN

TI Tricyclic nitrogen heterocycles as phosphodiesterase IV inhibitors  
GI

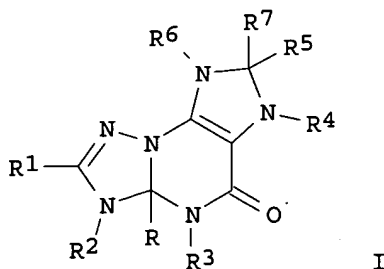


AB Tricyclic N heterocycles I [R1 = C1-5 alkyl, C5-6 cycloalkyl, Ph, PhCH2, 5- or 6-membered heterocyclic ring; R2 = C1-5 alkyl, C2-4 alkenyl; R3 = (substituted) C1-5 alkyl, (substituted) C5-6 cycloalkyl] and their salts are phosphodiesterase IV inhibitors and are potentially useful as vasodilators, inflammation inhibitors, and antiallergic agents. Thus, I (R1 = cyclopentyl, R2 = n-Pr, R3 = i-Pr) inhibited human monocyte phosphodiesterase IV with an IC50 of 0.018  $\mu$ m. A tablet formulation contained I 80, corn starch 190, lactose 55, microcryst. cellulose 35, PVP

15, Na carboxymethylstarch 23, and Mg stearate 2 mg.  
AN 2000:420941 CAPLUS <<LOGINID::20061109>>  
DN 133:53696  
TI Tricyclic nitrogen heterocycles as phosphodiesterase IV inhibitors  
IN Hoffmann, Matthias; Jung, Birgit; Kuefner-Muehl, Ulrike; Meade, Christopher John Montague  
PA Boehringer Ingelheim Pharma K.-G., Germany  
SO PCT Int. Appl., 17 pp.  
CODEN: PIXXD2  
DT Patent  
LA German  
FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2000035428	A2	20000622	WO 1999-EP9086	19991124
	WO 2000035428	A3	20000928		
	W: CA, JP, MX, US				
	RW: AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE				
	DE 19858331	A1	20000621	DE 1998-19858331	19981217
	CA 2345752	AA	20000622	CA 1999-2345752	19991124
	EP 1140098	A2	20011010	EP 1999-959324	19991124
	R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, FI				
	US 6417190	B1	20020709	US 1999-458789	19991210
PRAI	DE 1998-19858331	A	19981217		
	US 1999-127777P	P	19990405		
	WO 1999-EP9086	W	19991124		
OS	MARPAT 133:53696				

L13 ANSWER 4 OF 5 CAPLUS COPYRIGHT 2006 ACS on STN  
TI Preparation of imidazotriazolopyrimidines as adenosine receptor antagonists  
GI



AB Title compds. [I; R1 = H, alkyl, phenyl(alkyl), alkoxy carbonyl, etc.; R2 or R3 = H, alkyl, phenylalkyl, heterocyclyl(alkyl), etc.; R4 or R6 = H, (amino)alkyl, CH<sub>2</sub>Ph, etc.; R4R7 or R6R7 = bond; R5 = H, alkyl, phenyl(alkyl), etc.] were prepared Thus, 7-amino-2-[(4-methoxybenzyloxy)methyl]-s-triazolo[1,5-a]pyrimidine-5-one was converted in 10 steps to I (R2 = bond, R1 = CH<sub>2</sub>OPh, R3 = Et, R4 or R6 = H, R4R7 or R6R7 = bond, R5 = cyclopentyl). Data for biol. activity of I were given.  
AN 2000:161287 CAPLUS <<LOGINID::20061109>>  
DN 132:194388  
TI Preparation of imidazotriazolopyrimidines as adenosine receptor antagonists  
IN Kufner-muhl, Ulrike; Kummer, Werner; Pohl, Gerald; Gaida, Wolfram; Lehr,

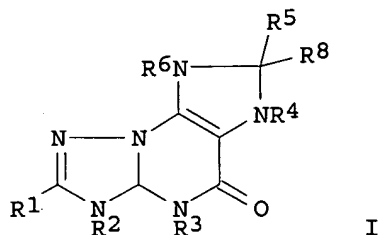
PA Erich; Mierau, Joachim; Weiser, Thomas; Carter, Adrian; Meade, Christopher  
 SO John Montague; Blech, Stefan; Hoffmann, Matthias  
 BOEHRINGER Ingelheim Pharma Kg, Germany; et al.  
 PCT Int. Appl., 77 pp.  
 CODEN: PIXXD2  
 DT Patent  
 LA German  
 FAN.CNT 3

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2000012511	A1	20000309	WO 1998-EP5455	19980827
	W: AU, BG, BR, BY, CA, CN, CZ, EE, HU, ID, IL, JP, KR, KZ, LT, LV, MX, NO, NZ, PL, RO, RU, SG, SI, SK, TR, UA, US, UZ, VN, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM RW: AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE				
	AU 9893474	A1	20000321	AU 1998-93474	19980827
	US 6492377	B1	20021210	US 2000-559806	20000426
PRAI	US 1998-90586P	P	19980625		
	US 1998-90587P	P	19980625		
	WO 1998-EP5455	A	19980827		
	US 1999-333408	A2	19990615		
	US 1999-333621	B2	19990615		

OS MARPAT 132:194388

RE.CNT 4 THERE ARE 4 CITED REFERENCES AVAILABLE FOR THIS RECORD  
 ALL CITATIONS AVAILABLE IN THE RE FORMAT

L13 ANSWER 5 OF 5 CAPLUS COPYRIGHT 2006 ACS on STN  
 TI Imidazotriazolopyrimidines as adenosine antagonists  
 GI



AB Imidazotriazolopyrimidines I [R1, R5 = (un)substituted alkyl, alkenyl, alkynyl, cycloalkyl, Ph, norbornyl, norbornenyl, adamantyl, noradamantyl, CO2H, CONH2, NH2, CHO; R2, R3 = (un)substituted alkyl; R2R7, R3R7, R4R8, R8R6 = bond; R4, R6 = H, alkyl, aminoalkyl, PhCH2; R2 and R3 or R4 and R6 cannot be present simultaneously] were prepared for use as adenosine antagonists. Thus, I [R1 = CH2OPh, R2R7, R4R8 = bond, R3 = Et, R5 = cyclopentyl, R4R8 = bond, II] was prepared from 4-MeOC6H4CH2OH, ClCH2CO2H, aminoguanidine cyclopentanecarbonyl chloride, and phenol in 12 steps. II had a KiA1 receptor binding activity of 3.6 nM.

AN 1999:811248 CAPLUS <<LOGINID::20061109>>

DN 132:35717

TI Imidazotriazolopyrimidines as adenosine antagonists

IN Blech, Stefan; Carter, Adrian; Gaida, Wolfram; Hoffmann, Matthias;  
 KuefnerMuehl, Ulrike; Meade, Christopher John Montague; Pohl, Gerald

PA Boehringer Ingelheim Pharma K.-G., Germany

SO PCT Int. Appl., 69 pp.

CODEN: PIXXD2

DT Patent  
LA German  
FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 9965912	A1	19991223	WO 1999-EP4017	19990611
	W: AU, BG, BR, BY, CA, CN, CZ, EE, HU, ID, IL, IN, JP, KR, KZ, LT, LV, MX, NO, NZ, PL, RO, RU, SG, SI, SK, TR, UA, US, UZ, VN, YU, ZA, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM				
	RW: AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE				
	DE 19826843	A1	19991223	DE 1998-19826843	19980616
	CA 2327395	AA	19991223	CA 1999-2327395	19990611
	AU 9945112	A1	20000105	AU 1999-45112	19990611
	EP 1087973	A1	20010404	EP 1999-927950	19990611
	EP 1087973	B1	20030108		
	R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, FI				
	JP 2002518396	T2	20020625	JP 2000-554737	19990611
	AT 230748	E	20030115	AT 1999-927950	19990611
	ES 2186369	T3	20030501	ES 1999-927950	19990611
PRAI	DE 1998-19826843	A	19980616		
	WO 1999-EP4017	W	19990611		

OS MARPAT 132:35717

RE.CNT 2 THERE ARE 2 CITED REFERENCES AVAILABLE FOR THIS RECORD  
ALL CITATIONS AVAILABLE IN THE RE FORMAT

=> s 16

L14 26 L6

=> d l14 1-26 ti

L14 ANSWER 1 OF 26 CAPLUS COPYRIGHT 2006 ACS on STN

TI New pharmaceutical compositions based on anticholinergics and PDE 5-inhibitors

L14 ANSWER 2 OF 26 CAPLUS COPYRIGHT 2006 ACS on STN

TI Pharmaceutical compositions based on anticholinergics and etiprednol

L14 ANSWER 3 OF 26 CAPLUS COPYRIGHT 2006 ACS on STN

TI Medicaments for the prevention or treatment of heart failure comprising administration of an anticholinergic

L14 ANSWER 4 OF 26 CAPLUS COPYRIGHT 2006 ACS on STN

TI Inhalable pharmaceutical compositions containing an anticholinergic, salmeterol, and a steroid

L14 ANSWER 5 OF 26 CAPLUS COPYRIGHT 2006 ACS on STN

TI Inhalable drugs containing a new anticholinergic, formoterol and a steroid

L14 ANSWER 6 OF 26 CAPLUS COPYRIGHT 2006 ACS on STN

TI Medicaments for the treatment of urinary tract disorders comprising anticholinergic agents

L14 ANSWER 7 OF 26 CAPLUS COPYRIGHT 2006 ACS on STN

TI Anticholinergic pharmaceuticals for the prevention or treatment of alveolar pneumonia

L14 ANSWER 8 OF 26 CAPLUS COPYRIGHT 2006 ACS on STN

TI Pharmaceutical compositions based on benzoic acid esters and soluble TNF receptor fusion proteins

L14 ANSWER 9 OF 26 CAPLUS COPYRIGHT 2006 ACS on STN

TI Aerosol formulation for inhalation containing an anticholinergic agent

L14 ANSWER 10 OF 26 CAPLUS COPYRIGHT 2006 ACS on STN  
 TI Medicament compositions comprising a heterocyclic compound and an anticholinergic

L14 ANSWER 11 OF 26 CAPLUS COPYRIGHT 2006 ACS on STN  
 TI Medicaments for inhalation comprising an anticholinergic and a betamimetic

L14 ANSWER 12 OF 26 CAPLUS COPYRIGHT 2006 ACS on STN  
 TI Powder formulations for inhalation containing an anticholinergic agent

L14 ANSWER 13 OF 26 CAPLUS COPYRIGHT 2006 ACS on STN  
 TI HFC solution aerosol formulations containing an anticholinergic tropane derivative

L14 ANSWER 14 OF 26 CAPLUS COPYRIGHT 2006 ACS on STN  
 TI Long-acting drug combinations for the treatment of respiratory tract diseases composed of an anticholinergic agent and a beta2-agonist

L14 ANSWER 15 OF 26 CAPLUS COPYRIGHT 2006 ACS on STN  
 TI Aerosol inhalant formulations containing a diphenylpropionic acid scopine ester-type anticholinergic agent

L14 ANSWER 16 OF 26 CAPLUS COPYRIGHT 2006 ACS on STN  
 TI Inhalants containing 2,2-diphenylpropionic acid scopine ester N-metho salts as anticholinergic agent in combination with corticosteroids and betamimetics

L14 ANSWER 17 OF 26 CAPLUS COPYRIGHT 2006 ACS on STN  
 TI Aerosol inhalant formulations containing a diphenylpropionic acid scopine ester-type anticholinergic agent

L14 ANSWER 18 OF 26 CAPLUS COPYRIGHT 2006 ACS on STN  
 TI Pharmaceutical compositions for the treatment of respiratory tract diseases comprising novel anticholinergic agents and inhibitors of EGFR-kinase

L14 ANSWER 19 OF 26 CAPLUS COPYRIGHT 2006 ACS on STN  
 TI Pharmaceutical compositions based on novel anticholinergics and p38 kinase inhibitors

L14 ANSWER 20 OF 26 CAPLUS COPYRIGHT 2006 ACS on STN  
 TI Pharmaceutical compositions comprising novel anticholinergic agents and NK1-receptor antagonists for the treatment of respiratory tract diseases

L14 ANSWER 21 OF 26 CAPLUS COPYRIGHT 2006 ACS on STN  
 TI Pharmaceutical compositions comprising anticholinergic agents and phosphodiesterase IV (PDE-IV) inhibitors for the treatment of respiratory diseases

L14 ANSWER 22 OF 26 CAPLUS COPYRIGHT 2006 ACS on STN  
 TI Medicaments containing betamimetic drugs and a novel anticholinesterase drug for treating respiratory tract diseases

L14 ANSWER 23 OF 26 CAPLUS COPYRIGHT 2006 ACS on STN  
 TI Pharmaceutical combinations containing heterocyclic compounds and scopine diphenyl propionate as anticholinergic agent

L14 ANSWER 24 OF 26 CAPLUS COPYRIGHT 2006 ACS on STN  
 TI Compositions containing steroids and a scopine diphenylpropionic acid derivative as anticholinesterase drug for the treatment of respiratory tract diseases

L14 ANSWER 25 OF 26 CAPLUS COPYRIGHT 2006 ACS on STN  
TI Method for producing scopine esters of diarylalkanoic acids

L14 ANSWER 26 OF 26 CAPLUS COPYRIGHT 2006 ACS on STN  
TI Procedures for the production of new anticholinergic alkaloids as well as for their use in medicines

=> s l14 not py>2002  
4558921 PY>2002  
L15 0 L14 NOT PY>2002

=> s l14 not py>2003  
3500279 PY>2003  
L16 0 L14 NOT PY>2003

=> s l14 not py>2004  
2357559 PY>2004  
L17 1 L14 NOT PY>2004

=> d l17 ti

L17 ANSWER 1 OF 1 CAPLUS COPYRIGHT 2006 ACS on STN  
TI Pharmaceutical combinations containing heterocyclic compounds and scopine diphenyl propionate as anticholinergic agent

=> d l17 ti abs bib

L17 ANSWER 1 OF 1 CAPLUS COPYRIGHT 2006 ACS on STN  
TI Pharmaceutical combinations containing heterocyclic compounds and scopine diphenyl propionate as anticholinergic agent  
AB The invention concerns pharmaceutical combinations that contain heterocyclic compds., especially benzofuran and benzopyran derivs., and scopine di-Ph propionate or its salts as an anticholinergic agent; the compns. are formulated as inhalants and are used for the treatment of respiratory tract diseases. Thus a microcapsule included ( $\mu$ g): scopine diphenylpropionate methobromide 200; heterocyclic compound 200; lactose 4600.

AN 2003:837039 CAPLUS <<LOGINID::20061109>>  
DN 139:328380

TI Pharmaceutical combinations containing heterocyclic compounds and scopine diphenyl propionate as anticholinergic agent

IN Banholzer, Rolf; Meade, Christopher John Montague; Meissner, Helmut; Morschhaeuser, Gerd; Pairret, Michel; Pieper, Michael P.; Pohl, Gerald; Reichl, Richard; Speck, Georg

PA Boehringer Ingelheim Pharma G.m.b.H. & Co. K.-G., Germany

SO PCT Int. Appl., 60 pp.  
CODEN: PIXXD2

DT Patent

LA German

FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
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PI	WO 2003087049	A2	20031023	WO 2003-EP3670	20030409
	WO 2003087049	A3	20040205		
	W:	AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NI, NO, NZ, OM, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW			
	RW:	GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES,			

	FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE, SI, SK, TR,			
	BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG			
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